



A comparison between Yfiler[®] and RM Y-STRs in United Arab Emirates population



R. Alghafri^{a,*}, S. Alhammadi^b, K. Amiri^b

^a General Department of Forensic Sciences and Criminology, Dubai Police General Head Quarters, Dubai, United Arab Emirates

^b Biology Department, College of Science, UAE University, Al Ain, United Arab Emirates

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ABSTRACT

Rapidly Mutating Y-STRs (RM Y-STR) were proven to have significantly higher resolution in worldwide populations when compared to commonly used Y-STR kits. In order to contribute to the research that has been conducted so far, RM Y-STRs were investigated in parallel with Yfiler[®] kit in 327 male individuals from United Arab Emirates population. Such population is considerably isolated where Yfiler[®] and Powerplex[®] Y haplotypes were found to be shared between distantly related as well as non-related male individuals. In the present study, a comparison between Y-STR markers included in Yfiler[®] kit (DYS456, DYS389I, DYS390, DYS389II, DYS458, DYS19, DYS385, DYS393, DYS391, DYS439, DYS635, DYS392, YGATAH4, DYS437, DYS438, DYS448) with RM Y-STRs (DYF387S1, DYF399S1, DYF403S1a/b, DYF404S1, DYS449, DYS518, DYS526a/b, DYS547, DYS570, DYS576, DYS612, DYS626, and DYS627) is conducted. RM Y-STRs were analysed using previously published methods. Forensic parameters were calculated for each set of markers, including discrimination capacity, haplotype diversity and gene diversity, using Arlequin v3.5 software.

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1. Introduction

Y chromosome STR profiling is a very common method which is used in forensic investigations, especially in sexual assault cases where male fraction of samples are hardly recovered. Yfiler[®] is one of the widely used multiplex assays in forensic laboratories. Recently a set of 13 markers were discovered which demonstrate significantly higher mutation rate than conventional markers, a feature which has been proven to be effective in discriminating non-relatives males populations as well as close male relatives [1]. Although some of these markers were incorporated in the newly released commercially available multiplex assays, it is still important to investigate the efficiency of these markers in different populations before adopting them for routine analysis. This study presents a parallel comparison between Yfiler[®] assay and RM Y-STR.

2. Materials and methods

2.1. Samples collection and extraction

Buccal swabs were collected from 327 individuals of national Arab's in the UAE population as part of Y chromosome polymorphism in UAE population study [2]. Briefly, DNA was extracted from samples using the Biorobot Universal System from Qiagen[®] with silica membrane matrix method. Quantification step was performed using Quantifiler[™] DNA Quantification Kit using a 7500 Real Time System (Applied Biosystems) following the manufacturer protocol.

2.2. Amplification and detection

Three multiplexes were used to analyse each sample following previously published methods in order to obtain the 13 markers of RM Y-STRs [1]. Each multiplex was performed in a volume of 10 μ l. Previously genotyped male positive controls along with negative controls were used during amplification to ensure quality of the analysis. PCR products were analyzed by capillary electrophoresis by an ABI 3500 Genetic Analyzer for fragment length determination of the products using POP-4[™] polymer and LIZ500[™] as internal size standard.

* Corresponding author.

E-mail address: r.alghafri@hotmail.co.uk (R. Alghafri).

Table 1
Forensic parameters for Yfiler® and RM Y-STRs in UAE Arab's populations.

Forensic parameters	Yfiler®	RM Y-STRs
Number of samples	327	327
Number of haplotypes	283	327
Number of unique haplotypes	253	327
Discrimination capacity (%)	89.40%	100%
Haplotypes diversity	0.98	1

2.3. Statistics analysis

Forensic parameters including discrimination capacity (DC), haplotype diversity (HD) and gene diversity (GD) were calculated using Arlequin v3.5 software [3].

3. Results and discussion

Among the 327 Arab's male individuals from the UAE population investigated, 327 distinguishable haplotypes were observed with RM Y-STR (Discrimination capacity of 100%) compared to 253 distinguishable haplotypes observed with Yfiler® (Discrimination capacity of 89.4%) shown in Table 1. This result represents a significant difference of 10% increase in resolution, between the two sets of markers (paired *t*-test, $t = -3.994$, $p = 0.001$). This result is in agreement with previous studies conducted using RM Y-STR on worldwide populations [1,4]. Extremely high resolution is explained by the high rate of mutation which is likely introducing new haplotypes in almost every generation and therefore increasing gene diversity (Table 2) (Fig. 1). Comparing haplotypes produced in this study with a previously published database of 12272 RM Y-STR haplotypes [4],

Table 2
Gene diversity for Yfiler® and RM Y-STRs in UAE Arab's populations.

Yfiler® Loci	Gene diversity	RM Y-STRs Loci	Gene diversity
DYS456	0.8405	DYF387S1	0.9212
DYS389I	0.5898	DYF399S1	0.9926
DYS390	0.8360	DYF403S1a	0.9812
DYS389II	0.8249	DYF403S1b	0.8979
DYS458	0.9262	DYF404S1	0.8257
DYS19	0.7451	DYS449	0.8718
DYS385a/b	0.9512	DYS518	0.8382
DYS393	0.7216	DYS526a	0.8267
DYS391	0.5906	DYS526b	0.8821
DYS439	0.7856	DYS547	0.7812
DYS635	0.8357	DYS570	0.7325
DYS392	0.4369	DYS576	0.7333
YGATAH4	0.7637	DYS612	0.8199
DYS437	0.4777	DYS626	0.6211
DYS438	0.7314	DYS627	0.8021
DYS448	0.6910		

there was no shared haplotypes between UAE Arab's populations and other populations, which also reflect the uniqueness of haplotypes across populations. These findings suggest that the set of 13 RM Y-STRs has great value in forensic investigations. Therefore, application of new generation multiplex assay which has incorporated some or all of these markers is highly recommended in forensic laboratories.

Conflict of interest

None.

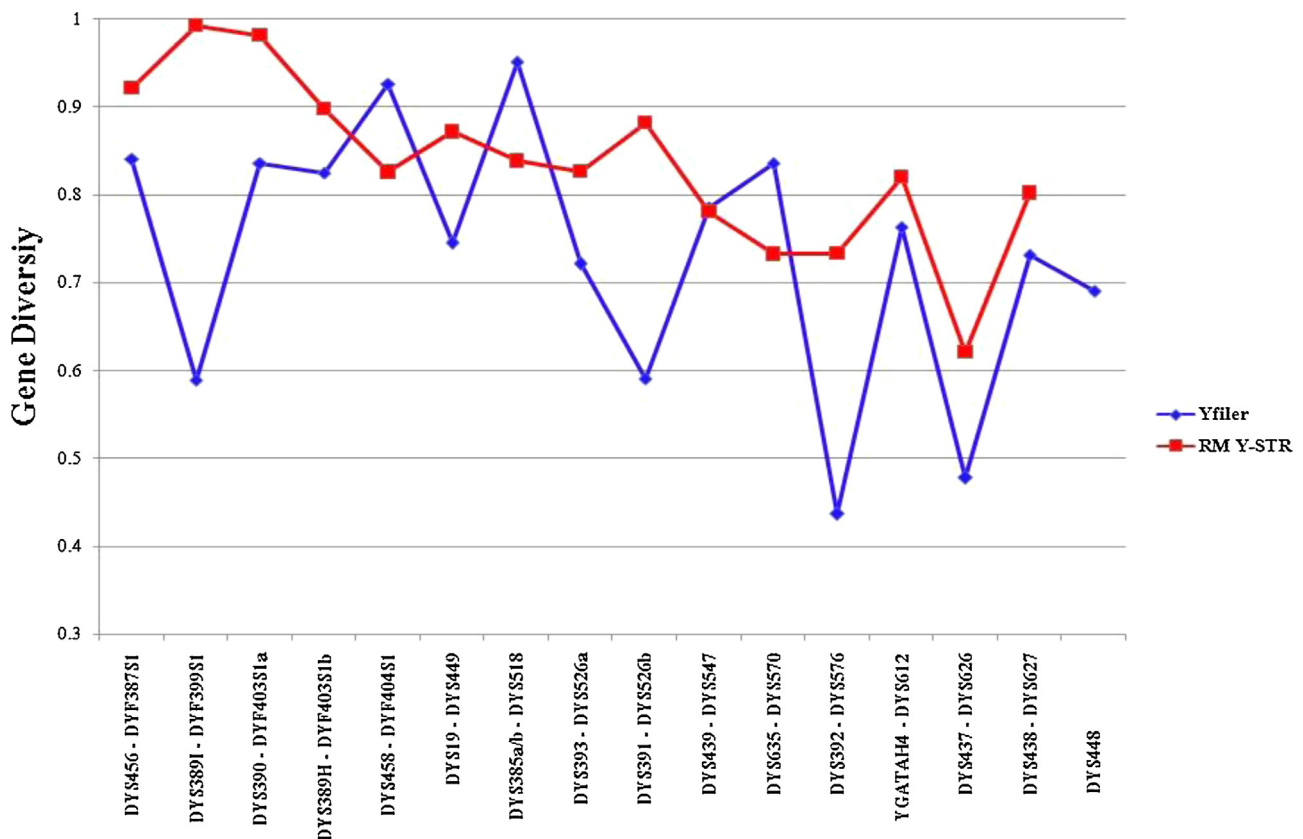


Fig. 1. Gene diversity for Yfiler® and RM Y-STRs in UAE Arab's population.

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